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531 Rec'd PCT/US 12 DEC 2001

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December 12, 2001

## BOX PCT

Commissioner for Patents  
Washington, D.C. 20231PCT/JP00/03877  
-filed June 15, 2000

Re: Application of Hiroyuki TANAKA, Masanori HASEGAWA, Tsuyoshi NOGUCHI, Katsuhiko HIGASHINO and Yosuke NISHIMURA  
CROSSLINKABLE ELASTOMER COMPOSITION AND MOLDED ARTICLE PRODUCED FROM SAME  
**Assignee: DAIKIN INDUSTRIES, LTD.**  
Our Ref: Q67530

Dear Sir:

The following documents and fees are submitted herewith in connection with the above application for the purpose of entering the National stage under 35 U.S.C. § 371 and in accordance with Chapter II of the Patent Cooperation Treaty:

- an executed Declaration and Power of Attorney.
- an English translation of the International Application.
- an English translation of Article 34 amendments
- an executed Assignment and PTO 1595 form.
- International Preliminary Examination Report
- International Search Report, Information Disclosure Statement, PTO Form 1449 with references.
- a Preliminary Amendment

It is assumed that copies of the Articles 19 and 34 amendments as required by § 371(c) will be supplied directly by the International Bureau, but if further copies are needed, the undersigned can easily provide them upon request.

The Government filing fee, after entry of the Article 34 Amendment and Preliminary Amendment, is calculated as follows:

Total claims	28	-	20	=	8	x	\$18.00	=	\$144.00
Independent claims	2	-	3	=		x	\$84.00	=	\$0.00
Base Fee									\$890.00

<b>TOTAL FILING FEE</b>	<u>\$1034.00</u>
<b>Recordation of Assignment</b>	<u>\$ 40.00</u>
<b>TOTAL FEE</b>	<u><u>\$1074.00</u></u>

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Checks for the statutory filing fee of \$1034.00 and Assignment recordation fee of \$40.00 are attached. You are also directed and authorized to charge or credit any difference or overpayment to Deposit Account No. 19-4880. The Commissioner is hereby authorized to charge any fees under 37 C.F.R. §§ 1.16, 1.17 and 1.492 which may be required during the entire pendency of the application to Deposit Account No. 19-4880. A duplicate copy of this transmittal letter is attached.

Priority is claimed from:

<u>Country</u>	<u>Application No</u>	<u>Filing Date</u>
Japan	174628/1999	June 21, 1999

Respectfully submitted,

A handwritten signature in black ink, appearing to read "AJR".

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Abraham J. Rosner  
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AJR/tmm  
Date: December 12, 2001

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PATENT APPLICATION

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re application of

HIROYUKI TANAKA et al.

Appln. No.: Not Yet Assigned

Confirmation No.: Not Yet Assigned

Group Art Unit: Not Assigned

Filed: December 20, 2001

Examiner: Not Assigned

For: CROSSLINKABLE ELASTOMER COMPOSITION AND MOLDED ARTICLE  
PRODUCED FROM SAME

**PRELIMINARY AMENDMENT**

Commissioner for Patents  
Washington, D.C. 20231

Sir:

Prior to examination, and after entry of the Article 34 Amendment, please amend the above-identified application as follows:

**IN THE SPECIFICATION:**

**Page 27, please delete the paragraph bridging pages 27 and 28, and replace it with  
the following new paragraph:**

A crosslinkable elastomer composition was prepared in the same manner as in Example 5 except that ultra fine powders of spherical quartz silica (1-FX available from Kabushiki Kaisha Tatsumori, specific surface area: 29.7 m<sup>2</sup>/g, average particle size: 0.38 µm) which had not been surface-treated with a silane coupling agent were used instead of the ultra fine powders of spherical synthetic quartz silica of Example 5 surface-treated with a silane coupling agent. Further the composition was molded into O-ring in the same manner as in Example 1. With

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Example 1. With respect to those composition and molded article, various characteristics thereof were measured in the same manner as in Example 1. The results are shown in Table 2.

**IN THE CLAIMS:**

**Please enter the following amended claims:**

3. (Amended) The crosslinkable elastomer composition of Claim 1, wherein an average particle size of the ultra fine powders of silicon oxide is from 0.01 to 0.05  $\mu\text{m}$ .
4. (Amended) The crosslinkable elastomer composition of Claim 1, wherein the ultra fine powders of silicon oxide are an amorphous silica.
5. (Amended) The crosslinkable elastomer composition of Claim 1, wherein the ultra fine powders of silicon oxide are surface-treated with hydrofluoric acid.
6. (Amended) The crosslinkable elastomer composition of Claim 1, wherein the ultra fine powders of silicon oxide are surface-treated with a silane coupling agent.
7. (Amended) The crosslinkable elastomer composition of Claim 1, wherein the ultra fine powders of silicon oxide are heat-treated at high temperature of not less than 400°C in an inert gas stream.
8. (Amended) The crosslinkable elastomer composition of Claim 1 which comprises 1 to 150 parts by weight of said ultra fine powders of silicon oxide on the basis of 100 parts by weight of the crosslinkable elastomer component.

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9. (Amended) The crosslinkable elastomer composition of Claim 1 which comprises 0.05 to 10 parts by weight of an organic peroxide, 0.1 to 10 parts by weight of a crosslinking aid and 1 to 150 parts by weight of said ultra fine powders of silicon oxide on the basis of 100 parts by weight of the crosslinkable elastomer component.

10. (Amended) The crosslinkable elastomer composition of Claim 1, wherein the crosslinkable elastomer component is a fluorine-containing elastomer.

12. (Amended) A molded article obtained by crosslinking and molding the crosslinkable elastomer composition of Claim 1.

14. (Amended) The molded article of Claim 12 which is used for semiconductor production apparatuses.

15. (Amended) The molded article of Claim 12 which is a sealing material to be used for sealing of semiconductor production apparatuses.

**Please add the following new claims 16-28:**

16. (New) The crosslinkable elastomer composition of Claim 2, wherein an average particle size of the ultra fine powders of silicon oxide is from 0.01 to 0.05  $\mu\text{m}$ .

17. (New) The crosslinkable elastomer composition of Claim 2, wherein the ultra fine powders of silicon oxide are an amorphous silica.

18. (New) The crosslinkable elastomer composition of Claim 2, wherein the ultra fine powders of silicon oxide are surface-treated with hydrofluoric acid.

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19. (New) The crosslinkable elastomer composition of Claim 2, wherein the ultra fine powders of silicon oxide are surface-treated with a silane coupling agent.

20. (New) The crosslinkable elastomer composition of Claim 2, wherein the ultra fine powders of silicon oxide are heat-treated at high temperature of not less than 400°C in an inert gas stream.

21. (New) The crosslinkable elastomer composition of Claim 2 which comprises 1 to 150 parts by weight of said ultra fine powders of silicon oxide on the basis of 100 parts by weight of the crosslinkable elastomer component.

22. (New) The crosslinkable elastomer composition of Claim 2 which comprises 0.05 to 10 parts by weight of an organic peroxide, 0.1 to 10 parts by weight of a crosslinking aid and 1 to 150 parts by weight of said ultra fine powders of silicon oxide on the basis of 100 parts by weight of the crosslinkable elastomer component.

23. (New) The crosslinkable elastomer composition of Claim 2, wherein the crosslinkable elastomer component is a fluorine-containing elastomer.

24. (New) The crosslinkable elastomer composition of Claim 23, wherein the fluorine-containing elastomer is a perfluoro elastomer.

25. (New) A molded article obtained by crosslinking and molding the crosslinkable elastomer composition of Claim 2.

26. (New) The molded article of Claim 25, wherein an amount of dioctylphthalate gas to be generated when heating at 200°C for 15 minutes is not more than 3 ppb.

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27. (New) the molded article of claim 25 which is used for semiconductor production apparatuses.

28. (New) The molded article of Claim 25 which is a sealing material to be used for sealing of semiconductor production apparatuses.

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REMARKS

Entry and consideration of this Amendment is respectfully requested.

Respectfully submitted,



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Date: December 12, 2001

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APPENDIX

VERSION WITH MARKINGS TO SHOW CHANGES MADE

IN THE SPECIFICATION:

The specification is changed as follows:

**Page 27, paragraph bridging pages 27 and 28:**

A crosslinkable elastomer composition was prepared in the same manner as in [Example 4] Example 5 except that ultra fine powders of spherical quartz silica (1-FX available from Kabushiki Kaisha Tatsumori, specific surface area: 29.7 m<sup>2</sup>/g, average particle size: 0.38 µm) which had not been surface-treated with a silane coupling agent were used instead of the ultra fine powders of spherical synthetic quartz silica of [Example 4] Example 5 surface-treated with a silane coupling agent. Further the composition was molded into O-ring in the same manner as in Example 1. With respect to those composition and molded article, various characteristics thereof were measured in the same manner as in Example 1. The results are shown in Table 2.

IN THE CLAIMS:

The claims are amended as follows:

3. (Amended) The crosslinkable elastomer composition of Claim 1 [or 2], wherein an average particle size of the ultra fine powders of silicon oxide is from 0.01 to 0.05 µm.

4. (Amended) The crosslinkable elastomer composition of [any of Claims 1 to 3] Claim 1, wherein the ultra fine powders of silicon oxide are an amorphous silica.

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5. (Amended) The crosslinkable elastomer composition of [any of Claims 1 to 3] Claim 1, wherein the ultra fine powders of silicon oxide are surface-treated with hydrofluoric acid.

6. (Amended) The crosslinkable elastomer composition of [any of Claims 1 to 3] Claim 1, wherein the ultra fine powders of silicon oxide are surface-treated with a silane coupling agent.

7. (Amended) The crosslinkable elastomer composition of [any of Claims 1 to 3] Claim 1, wherein the ultra fine powders of silicon oxide are heat-treated at high temperature of not less than 400°C in an inert gas stream.

8. (Amended) The crosslinkable elastomer composition of [any of Claims 1 to 7] Claim 1 which comprises 1 to 150 parts by weight of said ultra fine powders of silicon oxide on the basis of 100 parts by weight of the crosslinkable elastomer component.

9. (Amended) The crosslinkable elastomer composition of [any of Claims 1 to 8] Claim 1 which comprises 0.05 to 10 parts by weight of an organic peroxide, 0.1 to 10 parts by weight of a crosslinking aid and 1 to 150 parts by weight of said ultra fine powders of silicon oxide on the basis of 100 parts by weight of the crosslinkable elastomer component.

10. (Amended) The crosslinkable elastomer composition of [any of Claims 1 to 9] Claim 1, wherein the crosslinkable elastomer component is a fluorine-containing elastomer.

12. (Amended) A molded article obtained by crosslinking and molding the crosslinkable elastomer composition of [any of Claims 1 to 11] Claim 1.

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14. (Amended) The molded article of [Claim 12 or 13] Claim 12 which is used for semiconductor production apparatuses.

15. (Amended) The molded article of Claim [14] 12 which is a sealing material to be used for sealing of semiconductor production apparatuses.

**Claims 16-28 are added as new claims:**

## DESCRIPTION

CROSSLINKABLE ELASTOMER COMPOSITION  
AND MOLDED ARTICLE PRODUCED FROM SAME

5

## TECHNICAL FIELD

The present invention relates to a clean crosslinkable elastomer composition which contains ultra fine powders of silicon oxide and can provide a molded article to be used for semiconductor production apparatuses, for example, a sealing material, etc. used for sealing thereof, and relates to a molded article which is obtained from the composition, is excellent in mechanical strength and ensures a reduction in an amount of impurity gas to be released.

15

## BACKGROUND ART

In producing semiconductor elements, very high cleanliness is required, and the requirements for high cleanliness range over not only management of a production process of semiconductors but also a semiconductor production apparatus and parts thereof. If the parts of semiconductor production apparatuses are cleaned after built in the apparatuses, a degree of possible cleanliness is limited. Such parts are required to have been cleaned highly before built in the apparatuses. In production of semiconductors, there are various contaminations which come from parts of semiconductor production apparatuses, for example, so-called microparticles (impurity fine particles) generated or released from the parts and impurity gases (outgas) generated due to decomposition or modification of materials of the parts and mixed to an

atmosphere gas, which have an adverse effect particularly on accurate etching treatment of semiconductor elements.

Also cleanliness is required similarly in a molded article such as a sealing material for semiconductor production apparatuses which 5 the present invention can be particularly suitably applied to. The present applicants have attained high cleanliness of the sealing material itself by employing a special method of cleaning the sealing material after molding (JP-A-10-77781, JP-A-10-161988).

Such a sealing material is produced by crosslinking a 10 crosslinkable elastomer composition such as a rubber, and in order to improve mechanical properties such as compression set, there is a case where a metal oxide filler is added to the composition. There is a case where titanium oxide or silica (white carbon) is added as such a filler to enhance whiteness and improve plasma resistance (Japanese Patent 15 Nos. 2858198, 2783576).

In order to clean a molded article such as a sealing material, to which a metal oxide is added as a filler, particularly in order to eliminate impurity metals, the above-mentioned method of washing the sealing material after molding with an acid to extract impurity metals 20 and a method of adding a filler washed with an acid can be considered. In that case, it is necessary to crosslink an elastomer composition after the filler is added thereto. However when such a usual filler obtained by washing with an acid is used, peroxide crosslinking which has been employed commonly as a crosslinking method cannot be carried out. 25 Namely if peroxide crosslinking is intended to be conducted, in some cases, a crosslinking failure arises, which not only leads to lowering of mold-processability but also has an adverse effect on characteristics of a

molded article such as compression set, tensile strength and modulus.

Also in a production process of semiconductors, there is a case where a drying step such as plasma ashing is carried out. In that drying step, there is a case where microparticles (impurity fine particles) 5 arise from a molded article produced from an elastomer. Those microparticles as contamination sources also must be eliminated.

Particularly when silica is used, compression set is lowered. To improve it, the above-mentioned Japanese Patent No. 2783576 discloses use of ultra fine particles of white carbon having a pH value of 10 9 to 12 in 4 to 5 % by weight of aqueous dispersion.

Also in the above-mentioned Japanese Patent No. 2858198, a combination use of silica and carbon black is proposed to reduce an amount of microparticles to be generated in the drying process. The silica proposed therein is a dry silica having an average particle size of 1 15 to 50  $\mu\text{m}$ , preferably 10 to 40  $\mu\text{m}$ .

As a result of studies which have been made after the above-mentioned proposals, it was found that much amount of diethyl phthalate (DOP) are present in an outgas which is a problem. DOP has been used in a large amount, for example, as a plasticizer of plastics, 20 and is a compound which is present in a large amount in usual living environment.

In this regard, the composition disclosed in the Japanese Patent No. 2783576 has a problem that DOP is contained in an outgas in a high concentration.

25 Also the composition disclosed in the Japanese Patent No. 2858198 has an average particle size of as large as 1 to 50  $\mu\text{m}$ , and therefore when generation of microparticles arises, there is a problem

that in case of patterning of a standard line width of 0.3 to 0.35  $\mu\text{m}$  in 64 Mbit semiconductor, patterning defect occurs in some cases.

An object of the present invention is to provide a composition which is excellent in mechanical strength, particularly tensile strength  
5 even in case of a crosslinkable elastomer composition containing ultra fine powders of silicon oxide, and offers a reduced amount of outgas (particularly DOP and water) and microparticles.

#### DISCLOSURE OF INVENTION

10       Namely the present invention relates to the crosslinkable elastomer composition comprising a crosslinkable elastomer component and ultra fine powders of silicon oxide, in which the ultra fine powders of silicon oxide have, on surfaces thereof,  $100 \times 10^{19}$  or less, preferably  $90 \times 10^{19}$  or less hydroxyl groups per gram and have an average particle size  
15      of not more than 0.5  $\mu\text{m}$ , preferably from 0.01 to 0.03  $\mu\text{m}$ .

An amount of hydroxyl groups which are present on the surfaces of ultra fine powders of silicon oxide can be calculated from an amount of dehydration when heated at high temperature by the method mentioned hereinbelow.

20       In another aspect, the present invention relates to the crosslinkable elastomer composition comprising a crosslinkable elastomer component and ultra fine powders of silicon oxide, in which an adsorption amount of dioctyl phthalate on the ultra fine powders of silicon oxide is not more than 8  $\mu\text{g}/1\text{ g}$  of silicon oxide, preferably not  
25      more than 5  $\mu\text{g}/1\text{ g}$  of silicon oxide and an average particle size of the ultra fine powders of silicon oxide is not more than 0.5  $\mu\text{m}$ , preferably from 0.01 to 0.03  $\mu\text{m}$ .

It is preferable that the composition of the present invention contains 1 to 150 parts by weight (hereinafter referred to as "part") of the ultra fine powders of silicon oxide on the basis of 100 parts of the crosslinkable elastomer component. Also the composition can contain  
5 0.05 to 10 parts of an organic peroxide, 0.1 to 10 parts of a crosslinking aid and 1 to 150 parts of the ultra fine powders of silicon oxide on the basis of 100 parts of the crosslinkable elastomer component.

Examples of the preferred crosslinkable elastomer component are fluorine-containing elastomers, particularly perfluoro  
10 type elastomers.

The present invention also relates to the molded article obtained by crosslinking the above-mentioned crosslinkable elastomer composition, particularly molded articles to be used for semiconductor production apparatuses, such as a sealing material and the like to be  
15 used for sealing of semiconductor production apparatuses.

#### BEST MODE FOR CARRYING OUT THE INVENTION

The ultra fine powders of silicon oxide which are used in the present invention have  $100 \times 10^{19}$  or less, preferably  $90 \times 10^{19}$  or less,  
20 further preferably  $80 \times 10^{19}$  or less hydroxyl groups per gram on the surfaces thereof. When an amount of hydroxyl groups on the surfaces of ultra fine powders of silicon oxide is small, adsorption of water and DOP in the air can be reduced, and generation of steam and DOP which become an outgas can be decreased when the composition is processed  
25 into a molded article and used as parts of semiconductor production apparatuses.

As the ultra fine powders of silicon oxide which have a low

content of hydroxyl groups and a small amount of DOP adsorption, ultra fine powders of silica or ultra fine powders of synthetic quartz silica are used as they are or can be used after surface-treated with hydrofluoric acid or silane coupling agent.

5 Or ultra fine powders of silica or ultra fine powders of synthetic quartz silica are used after heat-treated at high temperature of not less than 400°C in an inert gas stream.

As the ultra fine powders of silica, there are dry silica, fumed silica and wet silica. In the present invention, dry silica or fumed silica is preferred from a point that an amount of outgas to be generated is small.

Examples of the ultra fine powders of synthetic quartz silica are, for instance, molten quartz silica, synthetic quartz silica, crystalline quartz silica, and the like.

15 The ultra fine powders of silica or synthetic quartz silica may  
be in either of irregular or spherical form.

The treatment with hydrofluoric acid may be carried out, for example, by dipping in 0.01 % dilute hydrofluoric acid for about 30 minutes, washing with ultra pure water and then drying in an inert gas containing no organic substance such as nitrogen gas.

Also the surface treatment with a silane coupling agent may be carried out through usual method. Examples of the preferred silane coupling agent are, for instance, vinyl silanes such as vinyltrimethoxysilane and vinyltriethoxysilane, amino silanes such as 25 N-(2-aminoethyl)-3-aminopropylmethyldimethoxysilane and N-(2-aminoethyl)-3-aminopropyltrimethoxysilane, 3-glycidoxypropyltrimethoxysilane, methacryloxypropyltrimethoxysilane,

3-mercaptopropyltrimethoxysilane, dimethyldichlorosilane, and the like. Particularly when peroxide crosslinking is carried out, treatment with a vinyl silane type silane coupling agent is preferred from the viewpoint of high reinforcing property and less releasing as microparticles.

5       The heat treatment at high temperature is carried out in an inert gas stream at a temperature of not less than 400°C, preferably 400° to 1,500°C, further preferably 500° to 1,200°C. By such heat treatment, thermal decomposition of hydroxyl groups being inherently present on surfaces of ultra fine powders of silica occurs and the number of  
10      hydroxyl groups are reduced remarkably and thus water adsorbing points can be decreased. As a result, an amount of water in the outgas can be reduced. At a temperature lower than 400°C, though it is possible to remove water already adsorbed, it is difficult to further carry out thermal decomposition of hydroxyl groups on the surfaces of ultra  
15      fine powders of silica. Heat treating time is from 30 to 240 minutes, usually from 30 to 180 minutes. A flow rate of the inert gas is not limited particularly, and may be usually from 0.1 to 20 liter/min. Examples of the inert gas are nitrogen gas, helium gas, argon gas, and the like.

20       An average particle size of the ultra fine powders of silicone oxide of the present invention is not more than 0.5 µm, preferably from 0.01 to 0.4 µm. Hitherto fine particles having a particle size of not less than 1 µm have been used as disclosed in the above-mentioned Japanese Patent No. 2858198. That is because as the particle size  
25      decreases, handling property of not only the ultra fine powders but also the crosslinkable composition is lowered. In the present invention, handling property (kneading property, etc.) of the crosslinkable

elastomer composition is not lowered.

Particularly when ultra fine powders having an average particle size of from about 0.01 µm to about 0.05 µm are used, even in case where a pattern of 0.2 µm distance between the patterning lines 5 (line width) is drawn on semiconductor element, no connection of lines occurs (no plugging occurs between the lines).

It is preferable to adjust a pH value of the ultra fine powders of silicon oxide to be acid or neutral (pH: not less than 3.0 and less than 9.0), particularly to be acid (pH: from 3.0 to 6.0) to reduce a density of 10 hydroxyl groups on the surfaces thereof. In that point, since the silica used for the composition disclosed in the above-mentioned Japanese Patent No. 2783576 has a pH value of as high as not less than 9 and has a lot of hydroxyl groups on the surfaces thereof, it can be considered that DOP is easily adsorbed to the silica and as a result of it, released into an 15 outgas in high concentration.

In case of application for molded articles for parts of semiconductor production apparatuses which require a very high cleanliness, it is preferable that a content of impurity metals other than silicon in the ultra fine powders of silicon oxide is adjusted to not more 20 than 10 ppm, preferably not more than 5 ppm.

Elastomer components are not limited particularly. When used as a material for producing a sealing material of semiconductor production apparatuses, a fluorine-containing elastomer and silicone elastomer are preferred.

25 Examples of the fluorine-containing elastomer are, for instance, those mentioned below.

A perfluoro elastomer comprising 40 to 90 % by mole of

tetrafluoroethylene, 10 to 60 % by mole of perfluorovinylether represented by the formula (1);



5

wherein  $R_f$  is a perfluoroalkyl group having 1 to 5 carbon atoms or a perfluoroalkyl(poly)ether group having 3 to 12 carbon atoms and 1 to 3 oxygen atoms, and 0 to 5 % by mole of monomer giving a curing site.

A vinylidene fluoride elastomer comprising 30 to 90 % by mole of  
10 vinylidene fluoride, 15 to 40 % by mole of hexafluoropropylene and 0 to 30 % by mole of tetrafluoroethylene.

A thermoplastic perfluoro elastomer which is a fluorine-containing multi-segment polymer comprising an elastomeric fluorine-containing polymer chain segment and a non-elastomeric fluorine-containing  
15 polymer chain segment, in which the elastomeric fluorine-containing polymer chain segment comprises 40 to 90 % by mole of tetrafluoroethylene, 10 to 60 % by mole of perfluorovinylether represented by the formula (1);

20



wherein  $R_f$  is a perfluoroalkyl group having 1 to 5 carbon atoms or a perfluoroalkyl(poly)ether group having 3 to 12 carbon atoms and 1 to 3 oxygen atoms, and 0 to 5 % by mole of monomer giving a curing site, and  
25 the non-elastomeric fluorine-containing polymer chain segment comprises 85 to 100 % by mole of tetrafluoroethylene and 0 to 15 % by mole of the formula (2);



wherein  $R_f^1$  is  $\text{CF}_3$  or  $\text{OR}_f^2$ , in which  $R_f^2$  is a perfluoroalkyl group having 1 to 5 carbon atoms.

- 5 A thermoplastic non-perfluoro elastomer which is a fluorine-containing multi-segment polymer comprising an elastomeric fluorine-containing polymer chain segment and a non-elastomeric fluorine-containing polymer chain segment, in which the elastomeric fluorine-containing polymer chain segment contains recurring units derived from 45 to 85 %
- 10 by mole of vinylidene fluoride and from at least one other monomer copolymerizable with vinylidene fluoride, respectively. Examples of the other monomer are hexafluoropropylene, tetrafluoroethylene, chlorotrifluoroethylene, trifluoroethylene, trifluoropropylene, tetrafluoropropylene, pentafluoropropylene, trifluorobutene,
- 15 tetrafluoroisobutene, perfluoro(alkyl vinyl ether), vinyl fluoride, ethylene, propylene, alkylvinylether, and the like.

A fluorine-containing elastomer having cold resistance which is obtained by radical polymerization in the presence of a diiodine compound and comprises 0.005 to 1.5 % by mole of fluorinated vinyl ether unit having iodine, 40 to 90 % by mole of vinylidene fluoride unit and 3 to 35 % by mole of perfluoro(methyl vinyl ether), in which as case demands, up to 25 % by mole of hexafluoropropylene unit and/or up to 40 % by mole of tetrafluoroethylene unit may be contained therein (JP-A-8-15753).

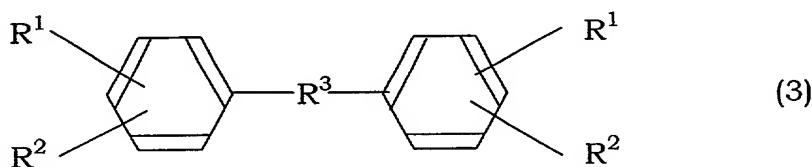
A copolymer of tetrafluoroethylene and propylene (USP 3467635).

- 25 As the silicone elastomer, preferred are, for example, a silicone rubber, fluoro silicone rubber, and the like.

The elastomer composition can be crosslinked and molded

into a desired form of product. Crosslinking is generally carried out by peroxide crosslinking. Also other known crosslinking methods are employed, for example, a triazine crosslinking system, in which a nitrile group is introduced as a crosslinking point into a fluorine-containing elastomer and a triazine ring is formed by using an organotin compound (for example, cf. JP-A-58-152041), an oxazole crosslinking system, in which a nitrile group is introduced similarly as a crosslinking point into a fluorine-containing elastomer and an oxazole ring is formed by using bisaminophenol (for example, cf. JP-A-59-109546), an imidazole crosslinking system, in which an imidazole ring is formed by using a tetraamine compound (for example, cf. JP-A-59-109546), a thiazole crosslinking system, in which a thiazole ring is formed by using bisaminothiophenol (for example, cf. JP-A-8-104789), and the like. Further radiation crosslinking method and electron beam crosslinking method may be employed.

Examples of a crosslinking agent used for oxazole crosslinking system, imidazole crosslinking system and thiazole crosslinking system are, for instance, a bisaminothiophenol crosslinking agent, bisaminophenol crosslinking agent and bisdiaminophenyl crosslinking agent represented by the formula (3):

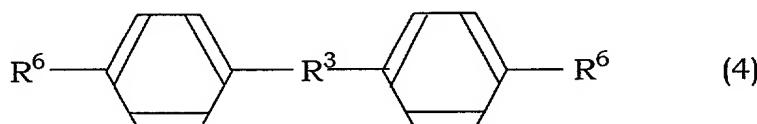


25

wherein R<sup>3</sup> is -SO<sub>2</sub>-, -O-, -CO-, an alkylene group having 1 to 6 carbon atoms, a perfluoroalkylene group having 1 to 10 carbon atoms or a

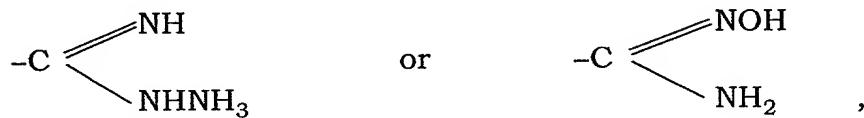
single bond, one of R<sup>1</sup> and R<sup>2</sup> is -NH<sub>2</sub> and another one is -NH<sub>2</sub>, -OH or -SH, preferably both of R<sup>1</sup> and R<sup>2</sup> are -NH<sub>2</sub>, bisamidorazone crosslinking agent represented by the formula (4):

5



in which R<sup>3</sup> is as defined above, R<sup>6</sup> is

10

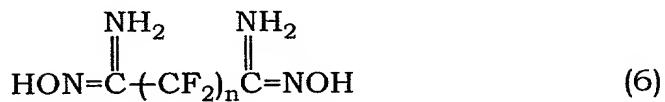


and bisamidoxime crosslinking agent represented by the formula (5) or

15 (6)



20 in which R<sub>f</sub><sup>3</sup> is a perfluoroalkylene group having 1 to 10 carbon atoms,

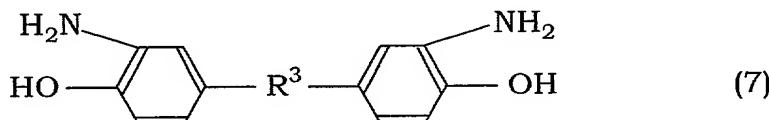


25 in which n is an integer of 1 to 10. Those bisaminophenol crosslinking agent, bisaminothiophenol crosslinking agent and bisdiaminophenyl crosslinking agent have been used for a crosslinking system wherein a

nitrile group is used as a crosslinking point. Further since they also react with carboxyl group and alkoxy carbonyl group, an oxazole ring, thiazole ring and imidazole ring are also formed in crosslinking systems having those functional groups as the crosslinking points and thus a 5 crosslinked product can be given.

Examples of the particularly preferred crosslinking agent are compounds having a plurality of 3-amino-4-hydroxyphenyl groups, 3-amino-4-mercaptophenyl groups or 3,4-diaminophenyl groups represented by the formula (7):

10



in which  $R^3$  is as defined above. Concrete examples thereof are, for 15 instance, 2,2-bis(3-amino-4-hydroxyphenyl)hexafluoropropane (general term: bis(aminophenol)AF), 2,2-bis(3-amino-4-mercaptophenyl)hexafluoropropane, tetraaminobenzene, bis-3,4-diaminophenylmethane, bis-3,4-diaminophenylether, 2,2-bis(3,4-diaminophenyl)hexafluoropropane, and the like.

20 An adding amount of the crosslinking agent is preferably from 0.1 to 10 parts on the basis of 100 parts of the elastomer.

When carrying out peroxide crosslinking, any of known organic peroxides which generate a peroxy radical under vulcanizing temperature condition may be used. Examples of the preferred organic 25 peroxide are di-t-butyl peroxide, dicumyl peroxide, 2,5-dimethyl-2,5-di(benzoylperoxy)hexane, 2,5-dimethyl-2,5-di(t-butylperoxy)hexane, 1,1-bis(t-butylperoxy)-3,5,5-trimethylcyclohexane, 2,5-dimethylhexane-

2,5-dihydroxy peroxide, t-butylcumyl peroxide,  $\alpha,\alpha'$ -bis(t-butylperoxy)-p-diisopropylbenzene, 2,5-dimethyl-2,5-di(t-butylperoxy)hexyne-3, benzoyl peroxide, t-butylperoxybenzene, t-butylperoxymaleate, t-butylperoxyisopropyl carbonate, and the like.

5 A content of the organic peroxide is usually from 0.05 to 10 parts, preferably from 1 to 5 parts on the basis of 100 parts of the fluorine-containing elastomer.

When the content of the organic peroxide is less than 0.05 part, the fluorine-containing elastomer is not crosslinked sufficiently, 10 and when more than 10 parts, physical properties of the crosslinked product are lowered.

In such peroxide crosslinking, a crosslinking aid such as a polyfunctional co-crosslinking agent can be used. As the polyfunctional co-crosslinking agent, those which are used together with 15 an organic peroxide in peroxide crosslinking of a fluorine-containing elastomer can be used. For example, there are bisolefins represented by triallylcyanurate, trimethallylisocyanurate, triallylisocyanurate, triallylformal, triallylphosphate, triallyltrimellitate, N,N'-m-phenylenebismaleimide, dipropargylterephthalate, diallylphthalate, 20 tetraallylterephthalamide, tris(diallylamine)-s-triazine, triallylphosphite, N,N-diallylacrylamide and 1,6-divinyldodecafluorohexane.

Also suitable are fluorine-containing triallylisocyanurate prepared by replacing a part of hydrogen atoms in three allyl groups of triallylisocyanurate with fluorine atoms having higher heat resistance, 25 and the like (cf. USP 4320216, WO98/00407, Klenovich, S.V. et al, Zh. Prikl. Khim. (Leningrad) (1987, 60(3), 656-8)).

A content of the crosslinking aid is usually from 0.1 to 10

parts, preferably from 0.5 to 5 parts on the basis of 100 parts of the fluorine-containing elastomer.

In addition, a processing aid, internal mold releasing agent, and the like may be added. Peroxide crosslinking can be carried out by 5 usual method, and no crosslinking failure recognized in conventional method arises.

The molded article of the present invention is cleaned significantly highly by treating, for example, through a special cleaning method disclosed in the above-mentioned JP-A-10-77781, namely a 10 method of washing with ultra-pure water, a method of washing with a clean organic compound in the form of liquid at a washing temperature or inorganic aqueous solution, a method of cleaning by dry etching or a method of extraction cleaning, and thus the molded article for semiconductor production apparatuses which ensures less releasing of 15 an outgas and is excellent in plasma resistance can be obtained.

The molded article of the present invention ensures less releasing of an outgas, and particularly generation of DOP when heating at 200°C for 15 minutes can be reduced to not more than 3 ppb, further not more than 1 ppb.

20 The crosslinkable elastomer composition of the present invention can be used suitably for a molded article for semiconductor production apparatuses, particularly for production of a sealing material for sealing of semiconductor production apparatuses, in which high cleanliness is demanded. Examples of the sealing material are O-ring, 25 square ring, gasket, packing, oil seal, bearing seal, lip seal, and the like.

In addition, the crosslinkable elastomer composition can be used for a variety of elastomer products, for example, diaphragm, tube,

hose, various rubber rolls, and the like. Also the composition can be used as a coating material and a lining material.

In the present invention, the semiconductor production apparatuses are not limited particularly to apparatuses for producing 5 semiconductors and encompass whole manufacturing equipment used in the field of semiconductors where a high degree of cleanliness is required, such as equipment for manufacturing a liquid crystal panel and plasma panel.

Examples of the semiconductor production apparatuses are 10 as follows.

(1) Etching system

Dry etching equipment

Plasma etching machine

Reactive ion etching machine

15 Reactive ion beam etching machine

Sputter etching machine

Ion beam etching machine

Wet etching equipment

Ashing equipment

20 (2) Cleaning system

Dry etching cleaning equipment

UV/O<sub>3</sub> cleaning machine

Ion beam cleaning machine

Laser beam cleaning machine

25 Plasma cleaning machine

Gas etching cleaning machine

Extractive cleaning equipment

Soxhlet extractive cleaning machine

High temperature high pressure extractive cleaning machine

Microwave extractive cleaning machine

Supercritical extractive cleaning machine

5 (3) Exposing system

Stepper

Coater and developer

(4) Polishing system

CMP equipment

10 (5) Film forming system

CVD equipment

Sputtering equipment

(6) Diffusion and ion implantation system

Oxidation and diffusion equipment

15 Ion implantation equipment

The present invention is then explained by means of examples, but is not limited to them.

#### EXAMPLE 1

20 The elastomer composition of the present invention was prepared by kneading 10 g of ultra fine powders of dry amorphous silica (REOLOSIL QS-10 available from Kabushiki Kaisha Tokuyama, specific surface area: 140 m<sup>2</sup>/g, average particle size: 0.02 µm), 1.0 g of PERHEXA 2.5B (available from NOF Corporation) and 3.0 g of triallylisocyanurate (TAIC) with 100 g of tetrafluoroethylene/perfluoro(alkyl vinyl ether) copolymer elastomer. The density of hydroxyl groups on the surfaces of ultra fine particles of

silicon oxide, DOP adsorption and contents of impurity metals were determined by the methods mentioned below. The results are shown in Table 1.

(Density of surface hydroxyl groups)

5       The ultra fine powders of silicon oxide are pre-treated by heating at 150°C for one hour to remove water absorbed in the air. Those pre-treated ultra fine particles of silicon oxide are measured by an amount of 5 g and further heated at 1,200° to 1,500°C until a weight reduction stops. Then an amount of the weight reduction is measured.

10       An amount of hydroxyl groups which are present on the surface (Density of surface hydroxyl groups: Number of hydroxyl groups/g) is calculated based on the results of measurement of weight reduction by the following equation.

15       Density of surface hydroxyl groups (Number of hydroxyl groups/g)

$$= (\text{Amount of weight reduction } (\mu\text{g/g}) \div (\text{Molecular weight of water} \times 10^6)) \times \text{Avogadro number} \times 2$$

(DOP adsorption)

20       Ultra fine powders of silicon oxide are added to DOP solution diluted with acetone, and after allowed to stand at room temperature for 15 minutes, the solution is heated at 60°C for 15 minutes to evaporate acetone. Thus the ultra fine powders of silicon oxide contaminated forcedly with DOP are prepared. Those ultra fine powders of silicon 25 oxide contaminated forcedly with DOP are put in a glass tube. After sealed, the tube is heated at 200°C for 15 minutes, and then generated gas is collected in a cooled trap tube. The trapped substance is rapidly

heated and analyzed with a gas chromatograph (GC-14A available from Shimadzu Corporation, Column: UA-15 available from Shimadzu Corporation). An amount of DOP released from the ultra fine particles of silicon oxide is calculated from a peak area of the obtained chart.

5 By using the results of the calculation, an amount of DOP which can be adsorbed by individual particles of ultra fine powders of silicon oxide is calculated (an amount of DOP adsorbed strongly on the ultra fine powders of silicon oxide).

Amount of DOP which can be adsorbed ( $\mu\text{g/g}$ )

10  $= \text{Amount of DOP forcedly adsorbed } (\mu\text{g/g}) - \text{Amount of released DOP } (\mu\text{g/g})$

(Weight reduction by heating at 200°C)

15 1.0 Gram of ultra fine powders of silicon oxide is put in an aluminum vessel and heated at 200°C until a change in weight reduction stops (usually about two hours). A weight reduction per unit surface area (% by weight/ $\text{m}^2$ ) by heating at 200°C is calculated from a change in weight after the heating by using the following equation.

20 Weight reduction per unit surface area by heating (% by weight/ $\text{m}^2$ )

$$= \frac{\text{Weight before heating (g)} - \text{Weight after heating (g)}}{\text{Weight before heating (g)}} \div \text{Specific surface area } (\text{m}^2/\text{g})$$

(Contents of impurity metals)

25 0.1 Gram of ultra fine powders of silicon oxide is put in a platinum crucible, and after diffused and melted in 5 ml of 50 % hydrofluoric acid in a hot bath, is diluted with ultra pure water.

Contents of metals of that solution are determined through atomic absorption analysis by using an atomic absorption photometer (Z8000 available from Hitachi, Ltd.). Metals intended to detect are Na, K and Cu. The metal content in the filler is determined by the following equation.

Metal content (ppm by weight)

$$= \frac{\text{Concentration in solution} \\ (\text{ppm by weight})}{\text{Weight of filler (g)}} \times \text{Weight of solution (g)}$$

10

The density of surface hydroxyl groups, DOP adsorption and contents of impurity metals of ultra fine powders of silicon oxide used in Examples 2 to 4 and Comparative Examples 1 to 2 are also shown in Table 1.

15

The ultra fine powders of silicon oxide in Table 1 are as follows.

REOLOSIL QS-10:

Ultra fine powders of amorphous silica available from  
Kabushiki Kaisha Tokuyama

20

REOLOSIL QS-40:

Ultra fine powders of amorphous silica available from  
Kabushiki Kaisha Tokuyama

REOLOSIL DM-10:

Ultra fine powders of amorphous silica available from  
Kabushiki Kaisha Tokuyama

25

Cab-O-Sil M-7D:

Ultra fine powders of fumed silica available from Cabot

- continued -

Contents of impurity  
metals (ppm by  
weight)

Na	0.5	0.02	2.0	2.0	0.07	0.3	0.2
K	0.05	0.01	0.1	0.1	0.01	0.04	0.03
Ca	0.1	0.08	0.5	0.5	0.02	0.2	0.04
Fe	0.3	0.3	0.2	0.2	0.1	0.3	0.05
Ni	0.005	0.0	0.09	0.09	0.0	0.02	0.0
Cu	0.0	0.0	0.01	0.01	0.003	0.0	0.0
Cr	0.02	0.01	0.04	0.04	0.0	0.05	0.0
Mg	0.007	0.01	0.01	0.01	0.0	0.01	0.002
Zn	0.07	0.06	0.1	0.1	0.03	0.06	0.05

TABLE 1

	Ex. 1	Ex. 2	Ex. 3	Ex. 4	Ex. 5	Com. Ex. 1	Com. Ex. 2
Ultra fine powders of silicon oxide	REOLOSIL QS-10	REOLOSIL DM-10	Cab-O-Sil M-7D	Cab-O-Sil M-7D (heat-treated)	1-FX treated with QS-40 vinyl silane	REOLOSIL 1-FX	
Density of surface hydroxyl groups (No. of groups/g)	$70 \times 10^{19}$	$6 \times 10^{19}$	$1 \times 10^{14}$	$1 \times 10^7$	$3 \times 10^{19}$	$190 \times 10^{19}$	$150 \times 10^{19}$
DOP absorption ( $\mu\text{g/g}$ )	1.9	0.8	0.0	0.0	0.6	14.2	8.1
Weight reduction by heating at $200^\circ\text{C}$ (% by weight/ $\text{m}^2$ )	5.0	4.0	3.0	0.1	4.0	10.0	9.0
Average particle size ( $\mu\text{m}$ )	0.02	0.03	0.02	0.02	0.38	0.01	0.38

- continued -

Specialty Chemicals, Inc.

1-FX:

Ultra fine powders of synthetic quartz silica available from  
Kabushiki Kaisha Tatsumori

The elastomer composition was subjected to press-crosslinking (primary crosslinking) at 160°C for 10 minutes and then crosslinking (secondary crosslinking) in an oven at 180°C for four hours to give an O-ring (AS-568A-214). Also a vulcanization curve of the 5 composition at 160°C was determined with JSR Curastometer Model II (available from Nichigo Shoji Kabushiki Kaisha), and a minimum viscosity (ML), degree of vulcanization (MH), induction time (T10) and optimum vulcanizing time (T90) were obtained. Further handling property of the composition was determined in the manner mentioned 10 below. The results are shown in Table 2.

(Processability)

Easiness of handling in the range of from kneading step to crosslinking step of the crosslinkable elastomer composition, for example, easy winding onto a roll and easy cutting at kneading and 15 flowability of the elastomer at crosslinking is synthetically evaluated with naked eyes.

Evaluation was made with naked eyes under the following criteria.

- A: No defects are found, and processing is carried out smoothly.
- 20 B: Almost no defect is found. Time and labor are required somewhat particularly at the kneading step.
- C: Much time and labor are required in the steps from the kneading step to the crosslinking step, but processing can be finished somehow.
- D: Processing cannot be carried out unless the processing conditions are 25 made strict considerably.

Mechanical properties and an amount of released DOP of the O-ring were measured as follows. The results are shown in Table 2.

(Mechanical properties)

Physical properties in normal state and compression set (200°C, 70 hours, compression by 25 %) were measured according to JIS K 6301.

5 (Amount of released DOP gas)

The O-ring (AS568A-214) is put in a glass tube. After sealed, the tube is heated at 200°C for 15 minutes, and then generated gas is collected in a cooled trap tube. The trapped substance is rapidly heated and analyzed with a gas chromatograph (GC-14A available from 10 Shimadzu Corporation, Column: UA-15 available from Shimadzu Corporation). An amount of DOP released from the O-ring is calculated from a peak area of the obtained chart.

(Amount of water generation)

An O-ring (AS568A-214) is put in a glass tube. After sealed, 15 the tube is heated at 200°C for 30 minutes, and then an amount of generated water is measured with Karl Fischer type water meter (available from Hiranuma Sangyo Kabushiki Kaisha).

## EXAMPLE 2

20 A crosslinkable elastomer composition was prepared in the same manner as in Example 1 except that ultra fine powders of dry amorphous silica (REOLOSIL DM-10 available from Kabushiki Kaisha Tokuyama, specific surface area: 120 m<sup>2</sup>/g, average particle size: 0.03 µm) surface-treated with monomethyltrichlorosilane (silane coupling agent) were used instead of the dry amorphous silica of Example 1. Further the composition was molded into O-ring in the same manner as 25 in Example 1. With respect to those composition and molded article,

various characteristics thereof were measured in the same manner as in Example 1. The results are shown in Table 2.

### EXAMPLE 3

5 A crosslinkable elastomer composition was prepared in the same manner as in Example 1 except that ultra fine powders of fumed silica (Cab-O-Sil M-7D (compression molding grade) available from Cabot Specialty Chemicals, Inc., specific surface area: 200 m<sup>2</sup>/g, average particle size: 0.02 µm) were used instead of the amorphous silica.

10 Further the composition was molded into O-ring in the same manner as in Example 1. With respect to those composition and molded article, various characteristics thereof were measured in the same manner as in Example 1. The results are shown in Table 2.

15

### EXAMPLE 4

A crosslinkable elastomer composition was prepared in the same manner as in Example 1 except that ultra fine powders of fumed silica (Cab-O-Sil M-7D (compression molding grade) available from Cabot Specialty Chemicals, Inc., specific surface area: 200 m<sup>2</sup>/g, average particle size: 0.02 µm) which had been subjected to heating at 600°C in a nitrogen stream for two hours and then allowing to stand for cooling down to room temperature in a nitrogen stream were used instead of the amorphous silica. Further the composition was molded into O-ring in the same manner as in Example 1. With respect to those composition and molded article, various characteristics thereof were measured in the same manner as in Example 1. The results are shown in Table 2.

quartz silica (1-FX available from Kabushiki Kaisha Tatsumori, specific surface area:  $29.7 \text{ m}^2/\text{g}$ , average particle size:  $0.38 \mu\text{m}$ ) which had not been surface-treated with a silane coupling agent were used instead of the ultra fine powders of spherical synthetic quartz silica of Example 4 5 surface-treated with a silane coupling agent. Further the composition was molded into O-ring in the same manner as in Example 1. With respect to those composition and molded article, various characteristics thereof were measured in the same manner as in Example 1. The results are shown in Table 2.

#### COMPARATIVE EXAMPLE 1

A crosslinkable elastomer composition was prepared in the same manner as in Example 1 except that ultra fine powders of dry amorphous silica having many hydroxyl groups on the surfaces thereof  
5 (REOLOSIL QS-40 available from Kabushiki Kaisha Tokuyama, specific surface area:  $380 \text{ m}^2/\text{g}$ , average particle size:  $0.01 \mu\text{m}$ ) were used as the dry amorphous silica. Further the composition was molded into O-ring in the same manner as in Example 1. With respect to those composition and molded article, various characteristics thereof were  
10 measured in the same manner as in Example 1. The results are shown in Table 2.

#### EXAMPLE 5

A crosslinkable elastomer composition was prepared in the same manner as in Example 1 except that ultra fine powders of spherical synthetic quartz silica (1-FX treated with vinyl silane available from Kabushiki Kaisha Tatsumori, specific surface area:  $29.7 \text{ m}^2/\text{g}$ , average particle size:  $0.38 \mu\text{m}$ ) surface-treated with a vinyl silane coupling agent were used instead of the dry amorphous silica. Further the composition was molded into O-ring in the same manner as in Example 1. With respect to those composition and molded article, various characteristics thereof were measured in the same manner as in Example 1. The results are shown in Table 2.

#### COMPARATIVE EXAMPLE 2

A crosslinkable elastomer composition was prepared in the same manner as in Example 4 except that ultra fine powders of spherical

TABLE 2

Components (part by weight)	Ex. 1	Ex. 2	Ex. 3	Ex. 4	Com. Ex. 1	Com. Ex. 5	Com. Ex. 2
Crosslinkable elastomer	100	100	100	100	100	100	100
Crosslinking agent	3	3	3	3	3	3	3
Crosslinking aid	1	1	1	1	1	1	1
Ultra fine powders of silicon oxide	10	10	10	10	10	10	10
Processability	A	A	A	A	B	B	B
Vulcanization characteristics							
ML (kg·f)	0.31	0.18	0.32	0.32	0.45	0.1	0.11
MH (kg·f)	8.4	8.2	8.15	8.15	8.85	6.25	5081
T10 (min)	0.6	0.5	0.5	0.5	0.5	0.7	0.7
T90 (min)	1.3	1.4	1.2	1.2	1	1.3	1.2

- continued -

- continued -

Physical properties in normal state

M100 (kg·f)	124	107	128	128	153	101	79
Tensile strength (kg·f)	220	212	233	233	213	182	170
Elongation (%)	135	148	138	138	127	125	155
Hardness (JIS A)	80	80	80	80	82	71	74
Compression set	17	16	11	11	17	12	12
Amount of released DOP (ppb)	0.92	0.41	0.00	0.00	7.00	0.31	3.95
Amount of generated water (ppm)	500	450	388	180	450	1000	900

## EXAMPLE 6

(Plasma resistance: Weight reduction)

The O-rings (AS-568A-214) produced, respectively in Examples 1 to 4, Comparative Example 1, Example 5 and Comparative Example 2 were put in petri dishes made of glass and then heated at 150°C for 60 minutes in nitrogen atmosphere to give samples.

Those samples were subjected to treating by plasma irradiation under the following conditions, and a weight reduction (% by weight) after the irradiation was measured to determine a change in weight. The results are shown in Table 3.

Plasma irradiation device used:

PX-1000 available from Kabushiki Kaisha Samco  
International Kenkyusho

Irradiation conditions:

Oxygen ( $O_2$ ) plasma irradiation treatment

Gas flow: 200 sccm

RF output: 400 W

Pressure: 300 mTorr

Etching time: 1 hr, 2 hr, 3 hr

Frequency: 13.56 MHz

$CF_4$  plasma irradiation treatment

Gas flow: 200 sccm

RF output: 400 W

Pressure: 300 mTorr

Etching time: 1 hr, 2 hr, 3 hr

Frequency: 13.56 MHz

Irradiation method:

In order to stabilize an atmosphere in a chamber of the plasma irradiation device, gas replacement is carried out for five minutes with empty chamber for pre-treatment of the chamber. Then the petri dish having the sample therein is placed at the center between the RF electrodes, and the irradiation is carried out under the above-mentioned conditions.

Measurement of weight:

A weight is measured up to two decimal places by using an electronic analyzing balance 2006MPE available from Sartorius GMBH and rounded to one decimal place.

Three samples are used for each kind and evaluation is made with an average thereof.

TABLE 3  
Weight reduction (% by weight)

	Oxygen plasma			CF <sub>4</sub> plasma		
	Irradiation time (hr)			Irradiation time (hr)		
	1	2	3	1	2	3
Com. Ex. 1	0.25	0.33	0.39	0.13	0.23	0.32
Ex. 1	0.21	0.28	0.33	0.12	0.21	0.30
Ex. 2	0.18	0.24	0.28	0.10	0.18	0.24
Ex. 3	0.21	0.29	0.36	0.12	0.23	0.29
Ex. 4	0.21	0.29	0.36	0.12	0.23	0.29
Com. Ex. 2	0.23	0.32	0.40	0.12	0.18	0.26
Ex. 5	0.22	0.33	0.41	0.10	0.17	0.24

#### EXAMPLE 7

(Plasma resistance: Number of microparticles generated)

Plasma resistance (Number of microparticles generated) of the O-rings (AS-568A-214) produced, respectively in Examples 1 to 4, 5 Comparative Example 1, Example 5 and Comparative Example 2 was determined by the method mentioned below. The results are shown in Table 4.

(Plasma resistance test: Number of microparticles generated)

Oxygen plasma or  $\text{CF}_4$  plasma was generated under the 10 conditions of a vacuum pressure of 50 mTorr, an oxygen flow or  $\text{CF}_4$  flow of 200 cc/min, electric power of 400 W and a frequency of 13.56 MHz by using Plasma Dry Cleaner Model PX-1000 available from Kabushiki Kaisha Samco International Kenkyusho, and the generated oxygen plasma or  $\text{CF}_4$  plasma was irradiated to the samples (O-ring) under the 15 reactive ion etching (RIE) conditions for three hours. After the irradiation, a supersonic wave was applied to the samples in ultra pure water at 25°C for one hour to take out free microparticles in water, and then the number of microparticles (per liter) having a particle size of not less than 0.2  $\mu\text{m}$  was measured by a fine particle meter method (a 20 method of emitting light to ultra pure water containing microparticles which was flowed into a sensor part and then electrically measuring amounts of transmitted light and scattered light with a submerged microparticle counter). In Table 4, values converted to the number of 25 microparticles per unit area ( $\text{cm}^2$ ) of the O-ring are also shown in the parentheses.

TABLE 4  
Number of microparticles generated ( $\times 10^4$ /liter)

Sample	Before irradiation	After one hour irradiation of oxygen plasma	After one hour irradiation of CF <sub>4</sub> plasma
Com. Ex. 1	7.6	32.9 (25.3)	11.3 (3.7)
Ex. 1	5.2	30.9 (25.7)	21.7 (16.5)
Ex. 2	8.4	24.6 (16.2)	46.3 (37.9)
Ex. 3	8.6	13.2 (4.6)	8.5 (0)
Ex. 4	8.6	13.2 (4.6)	8.5 (0)
Com. Ex. 2	5.4	47.2 (41.8)	48.5 (43.1)
Ex. 5	5.4	47.7 (42.3)	29.3 (23.9)

#### INDUSTRIAL APPLICABILITY

The crosslinkable elastomer composition of the present invention which comprises ultra fine powders of silicon oxide having a small density of surface hydroxyl groups and less adsorbing DOP is  
5 excellent in mechanical properties, and an amount of outgas is small and thus the composition is suitable as a very clean material for molded article for semiconductor production apparatuses.

ATT 34 AMDT

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## CLAIMS

1. (Amended) A crosslinkable elastomer composition comprising a crosslinkable elastomer (excluding a fluoro silicone rubber) component and ultra fine powders of silicon oxide; said ultra fine powders of silicon oxide have  $100 \times 10^{19}$  or less hydroxyl groups per gram on surfaces thereof and an average particle size of not more than 0.5  $\mu\text{m}$ .

2. (Amended) A crosslinkable elastomer composition comprising a crosslinkable elastomer (excluding a fluoro silicone rubber) component and ultra fine powders of silicon oxide; said ultra fine powders of silicon oxide have dioctylphthalate adsorption of not more than 8  $\mu\text{g}$  per 1 g of silicon oxide and an average particle size of not more than 0.5  $\mu\text{m}$ .

3. The crosslinkable elastomer composition of Claim 1 or 2, wherein an average particle size of the ultra fine powders of silicon oxide is from 0.01 to 0.05  $\mu\text{m}$ .

4. The crosslinkable elastomer composition of any of Claims 1 to 3, wherein the ultra fine powders of silicon oxide are an amorphous silica.

5. The crosslinkable elastomer composition of any of Claims 1 to 3, wherein the ultra fine powders of silicon oxide are surface-treated with hydrofluoric acid.

6. The crosslinkable elastomer composition of any of Claims

1 to 3, wherein the ultra fine powders of silicon oxide are surface-treated with a silane coupling agent.

7. The crosslinkable elastomer composition of any of Claims  
5 1 to 3, wherein the ultra fine powders of silicon oxide are heat-treated at high temperature of not less than 400°C in an inert gas stream.

8. The crosslinkable elastomer composition of any of Claims  
10 1 to 7 which comprises 1 to 150 parts by weight of said ultra fine powders of silicon oxide on the basis of 100 parts by weight of the crosslinkable elastomer component.

9. The crosslinkable elastomer composition of any of Claims  
15 1 to 8 which comprises 0.05 to 10 parts by weight of an organic peroxide, 0.1 to 10 parts by weight of a crosslinking aid and 1 to 150 parts by weight of said ultra fine powders of silicon oxide on the basis of 100 parts by weight of the crosslinkable elastomer component.

10. The crosslinkable elastomer composition of any of  
20 Claims 1 to 9, wherein the crosslinkable elastomer component is a fluorine-containing elastomer.

11. The crosslinkable elastomer composition of Claim 10,  
wherein the fluorine-containing elastomer is a perfluoro elastomer.

12. A molded article obtained by crosslinking and molding the crosslinkable elastomer composition of any of Claims 1 to 11.

13. The molded article of Claim 12, wherein an amount of dioctylphthalate gas to be generated when heating at 200°C for 15 minutes is not more than 3 ppb.

5

14. The molded article of Claim 12 or 13 which is used for semiconductor production apparatuses.

10 15. The molded article of Claim 14 which is a sealing material to be used for sealing of semiconductor production apparatuses.

# Declaration and Power of Attorney for Patent Application

特許出願宣言書及び委任状

## Japanese Language Declaration

### 日本語宣言書

下記の氏名の発明者として、私は以下通り宣言します。

私の住所、私書箱、国籍は下記の私の氏名の後に記載された通りです。

下記の名称の発明に関して請求範囲に記載され、特許出願している発明内容について、私が最初かつ唯一の発明者（下記の氏名が一つの場合）もしくは最初かつ共同発明者であると（下記の名称が複数の場合）信じています。

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated next to my name.

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled

CROSSLINKABLE ELASTOMER COMPOSITION  
AND MOLDED ARTICLE PRODUCED FROM SAME

上記発明の明細書（下記の欄でx印がついていない場合は、本書に添付）は、

\_\_\_月\_\_\_日に提出され、米国出願番号または特許協定条約  
国際出願番号を \_\_\_\_\_ とし。  
(該当する場合) \_\_\_\_\_ に訂正されました。

the specification of which is attached hereto unless the following box is checked:

was filed on June 15, 2000  
as U.S. Application No. or PCT International  
Application No. PCT/JP00/03877  
and was amended on \_\_\_\_\_  
(if applicable)

私は、特許請求範囲を含む上記訂正後の明細書を検討し、内容を理解していることをここに表明します。

私は、連邦規則法典第37編第1条56項に定義されるとおり、特許資格の有無について重要な情報を開示する義務があることを認めます。

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to patentability as defined in Title 37, Code of Federal Regulations, § 1.56.

# Japanese Language Declaration

## 日本語宣言

私は、米国法典第35編119条(a)-(d)項又は365条(b)項に基づき下記の、米国以外の国の少なくとも一ヵ国を指定している特許協力条約365(a)項に基づく国際出願、又は外国での特許出願もしくは発明者証の出願についての外国優先権をここに主張するとともに、優先権を主張している。本出願の前に出願された特許または発明者証の外国出願を以下に、枠内をマークすることで、示しています。

### Prior foreign application(s) 外国での先行出願

174628/1999 (Number) (番号)	Japan (Country) (国名)
_____ (Number) (番号)	_____ (Country) (国名)
_____ (Number) (番号)	_____ (Country) (国名)

私たる、第35編米国法典119条(e)項に基いて下記の米国特許出願規定に記載された権利をここに主張いたします。

(Application No.) (出願番号)	(Filing Date) (出願日)
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私は、下記の米国法典第35編120条に基いて下記の米国特許出願に記載された権利、又は米国を指定している特許協力条約365条(c)に基づく権利をここに主張します。また、本出願の各請求範囲の内容が米国法典第35編112条第1項又は特許協力条約で規定された方法で先行する米国特許出願に開示されていない限り、その先行米国出願書提出日以降で本出願書の日本国内または特許協力条約国際提出日までの期間中に入手された、連邦規則法典第37編1条56項で定義された特許資格の有無に関する重要な情報について開示義務があることを認識しています。

(Application No.) (出願番号)	(Filing Date) (出願日)
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私は、私自身の知識に基づいて本宣言書中で私が行なう表明が真実であり、かつ私の入手した情報と私の信じるところに基づく表明が全て真実であると信じていること、さらに故意になされた虚偽の表明及びそれと同等の行為は米国法典第18編第1001条に基づき、罰金または拘禁、もしくはその両方により処罰されること、そしてそのような故意による虚偽の声明を行なえば、出願した、又は既に許可された特許の有効性が失われることを認識し、よってここに上記のごとく宣誓を致します。

I hereby claim foreign priority under Title 35, United States Code, § 119(a)-(d) or § 365 (b) of any foreign application(s) for patent or inventor's certificate, or § 365(a) of any PCT International application which designated at least one country other than the United States, listed below and have also identified below, by checking the box, any foreign application for patent or inventor's certificate, or PCT International application having a filing date before that of the application on which priority is claimed.

### Priority Claimed 優先権の主張

21/6/1999 (Day/Month/Year Filed) (出願年月日)	<input checked="" type="checkbox"/> <input type="checkbox"/> Yes No ありなし
_____ (Day/Month/Year Filed) (出願年月日)	<input type="checkbox"/> <input checked="" type="checkbox"/> Yes No ありなし
_____ (Day/Month/Year Filed) (出願年月日)	<input type="checkbox"/> <input checked="" type="checkbox"/> Yes No ありなし

I hereby claim the benefit under Title 35, United States Code, § 119(e) of any United States provisional application(s) listed below.

(Application No.) (出願番号)	(Filing Date) (出願日)
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I hereby claim the benefit under Title 35, United States Code, § 120 of any United States application(s), or § 365(c) of any PCT International application designating the United States, listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States or PCT International application in the manner provided by the first paragraph of Title 35, United States Code, § 112, I acknowledge the duty to disclose information which is material to patentability as defined in Title 37, Code of Federal Regulations, § 1.56 which became available between the filing date of the prior application and the national or PCT International filing date of this application.

(Status)(patented, pending, abandoned)  
(現況：特許許可済、係属中、放棄済)

(Status)(patented, pending, abandoned)  
(現況：特許許可済、係属中、放棄済)

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under Section 1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

# Japanese Language Declaration

## 日本語宣言

委任状： 私は下記の発明者として、本出願に関する一切の手続きを米特許商標局に対して遂行する弁理士または代理人として、下記の者を指名いたします。（弁護士、または代理人の氏名及び登録番号を明記のこと）

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発明者の署名	Inventor's signature	Date
住所	<u>Hiroyuki Tanaka</u> 16/10/2001	
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私書箱	Citizenship Japanese Post Office Address c/o Yodogawa-seisakusho, DAIKIN INDUSTRIES, LTD., 1-1, Nishihitotsuya, Settsu-shi, OSAKA 566-8585 JAPAN	
第二共同発明者	Full name of second joint inventor, if any <u>Masanori HASEGAWA</u>	
第二共同発明者	Second inventor's signature	Date
住所	<u>Masanori Hasegawa</u> 16/10/2001	
国籍	Residence <u>Settsu-shi, OSAKA 566-8585 JAPAN</u>	
私書箱	Citizenship Japanese Post Office Address c/o Yodogawa-seisakusho, DAIKIN INDUSTRIES, LTD., 1-1, Nishihitotsuya, Settsu-shi, OSAKA 566-8585 JAPAN	

（第三以降の共同発明者についても同様に記載し、署名すること）

(Supply similar information and signature for third and subsequent joint inventors.)

# Japanese Language Declaration

日本語宣言

第三の共同発明者の氏名（該当する場合）		Full name of third joint inventor, if any Tsuyoshi NOGUCHI
同第三発明者の署名	日付	Third inventor's signature Date <i>Tsuyoshi Noguchi</i> 16/10/2001
住所		Residence: Settsu-shi, OSAKA 566-8585 JAPAN
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第四の共同発明者の氏名（該当する場合）		Full name of fourth joint inventor, if any Katsuhiko HIGASHINO
同第四発明者の署名	日付	Fourth inventor's signature Date <i>Katsuhiko Higashino</i> 16/10/2001
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第五の共同発明者の氏名（該当する場合）		Full name of fifth joint inventor, if any Yosuke NISHIMURA
同第五発明者の署名	日付	Fifth inventor's signature Date <i>Yosuke Nishimura</i> 16/10/2001
住所		Residence Settsu-shi, OSAKA 566-8585 JAPAN
国籍		Citizenship Japanese
郵便の宛先		Post office address c/o Yodogawa-seisakusho, DAIKIN INDUSTRIES, LTD., 1-1, Nishihitotsuya, Settsu-shi, OSAKA 566-8585 JAPAN
第六の共同発明者の氏名（該当する場合）		Full name of sixth joint inventor, if any
同第六発明者の署名	日付	Sixth inventor's signature Date
住所		Residence
国籍		Citizenship
郵便の宛先		Post office address